AUG 23 2006

K062145

5. 510(k) Summary

General Information

Date Compiled	July 26, 2006
Classification	Class II
Trade Name	Modified Acumen Lead Delivery Sheath Models: LDS-10-57, LDS-10-64, LDS-10-57-90, and LDS-10-64-90
Submitter	Acumen Medical, Inc. 275 Santa Ana Court Sunnyvale, CA 94085 Tel: 408-530-1810 Fax: 408-530-1811
Contact	Marybeth Gamber Director, Regulatory Affairs

Intended Use

The Modified Acumen Lead Delivery Sheath is indicated for the introduction of various types of pacing or defibrillator leads and catheters..

Predicate Devices

Acumen Lead Delivery Sheath K051515
Manufactured by Acumen Medical, Inc.

Device Description

The Acumen Lead Delivery Sheath (LDS) is a single-use percutaneous catheter intended to introduce various types of pacing or defibrillator leads and catheters. The Modified Acumen LDS has a guidewire lumen and a lumen for introduction of leads.

Materials

All materials used in the manufacture of the Modified Acumen LDS are suitable for this use and have been used in numerous previously cleared products.

Testing

Appropriate testing has been performed and all components, subassemblies, and/or full devices met the required specifications for the completed tests.

Summary of Substantial Equivalence

Acumen Medical, Inc. believes the Modified Acumen Lead Delivery Sheath is substantially equivalent to the predicate product. The intended use, method of operation, methods of construction and materials used, are either identical or substantially equivalent to existing legally marketed predicate product.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

AUG 2 3 2006

Acumen Medical, Inc.
c/o Ms. Marybeth Gamber
Director, Regulatory Affairs
275 Santa Ana Court
Sunnyvale, CA 94085

Re: K062145
Trade/Device Name: Modified Acumen Lead Delivery Sheath
Regulation Number: 21 CFR §870.1340
Regulation Name: Catheter Introducer
Regulatory Class: Class II
Product Code: DYB
Dated: July 26, 2006
Received: July 27, 2006

Dear Ms. Marybeth Gamber:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.
This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (240) 276-0120. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,

B Zuckerman for

Bram D. Zuckerman, M.D.
Director
Division of Cardiovascular Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

4. Indications for Use Statement

510(k) Number (if known):	This application
Device Name:	Modified Acumen Lead Delivery Sheath
Indications for Use:	The Acumen Lead Delivery Sheath is indicated for the introduction of various types of pacing or defibrillator leads and catheters.

Prescription Use _X_ (Per 21 CFR 801 Subpart D) AND/OR Over-The-Counter Use_____ (Per 21 CFR 801 Subpart C)

PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of Cardiovascular Devices
510(k) Number K062145